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| APPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. |
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| 10/802,143 | 03/16/2004 | Toshihisa Takeyama | KON-1860 | 3282 |
| 20311 | 7590 | 05/24/2007 | | |
| LUCAS & MERCANTI, LLP 475 PARK AVENUE SOUTH 15TH FLOOR NEW YORK, NY 10016 | | | EXAMINER ANGEBRANNDT, MARTIN J | |
| | | | ART UNIT 1756 | PAPER NUMBER |
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary

Application No.

10/802,143

Applicant(s)

TAKEYAMA, TOSHIHISA

Examiner

Martin J. Angebrannt

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-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 3/20/07 & 3/9/07.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-10 and 12-20 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-10 and 12-20 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
- Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
 2. ☐ Certified copies of the priority documents have been received in Application No. _____.
 3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- 1) ☐ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☒ Information Disclosure Statement(s) (PTO/SB/08)
Paper No(s)/Mail Date 3/20/07.
- 4) ☐ Interview Summary (PTO-413)
Paper No(s)/Mail Date. _____.
- 5) ☐ Notice of Informal Patent Application
- 6) ☐ Other: _____.

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1. The response of the applicant has been read and given careful consideration. Priority has been perfected and the applicant is accorded the date of 03/24/2003. This removes the Sasa et al. WO 2004/017141 reference. Rejection of the previous office action not repeated below are withdrawn based upon the amendment to the claims.

2. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

3. Claims 1-4,6-9,14 and 16-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Korishima et al. JP 05-046061, in view of Dhar et al. '551.

Korishima et al. JP 05-046061 teaches composition for forming volume holograms where a dye cation [0007] and a borate anion [0008-0010] are combined with a free readical polymerizable systems using ethylene based compounds such as acrylic or acrylate monomers [0012]. Example 1 uses a cyanine dyes with a triphenyl-butyl boarate anoin combinaed with N,N-dimethylaniline, polyvinylbutyral, vinyl carboazole and 2-phenoxy acrylate coated on a glass plate. These were exposed using an 830 nm laser forming the interferences fringes and then finally cured using a halogen lamp thereby forming a reflection hologram. [00189-0022]. The sensitization is different from that of other photocurable composition, being much improved in the red and infrared. [0003-0004,0023].

Dhar et al. '551 teach in example 1, an acrylate monomer and CGI-784 as the photoinitiator mixed with matrix precursors dibutyltin dilaurate, diisocyanate terminated polypropylene glycol and dihydroxypolypropylene glycol, which are heated (13/65-15).

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Examples 3 and 4 are similar, place the composition between two glass slides with a spacer and after curing of the matrix are used to record holograms. The ability to form thick recording layers of more than 200 microns is disclosed. (3/13-19,4/3-12). Useful photoactive monomers including acrylates are disclosed as useful in this system. (6/51-67). A reduction in shrinkages of the hologram is also realized (7/1-48). The use of various reactions including hydrosilation and the like to form the matrix is disclosed. (6/26-50)

It would have been obvious to one skilled in the art to modify the cited example of Korishima et al. JP 05-046061 by adding the components to form an insitu matrix and place the composition between two glass slides with a spacer of 200-500 microns to allow a thicker hologram to be formed and reduce shrinkage as discussed by Dhar et al. '551.

The applicant apparently has interpreted the references as the N,N-dimethylaniline being the counter ion. This is not the case. Even casual review of formula (2) on page 4 of Korishima et al. JP 05-046061 shows the cyanine dye to be the counter ion. The examiner notes that cyanine dyes are specifically disclosed and claimed by the applicant. The positive charge (making it cationic) is shown on the nitrogen of the right pyridine ring of the cyanine dye. The applicants analysis is flawed on its face. The applicant argues that there is no motivation to use the dye/borate of the claims in the secondary references. The examiner holds that the modification asserted in the rejection is to add the precursors to allow the formation of the matrix and the thicker holographic recording medium, and starts from a reference using the dye/borate. There is a reasonable expectation of success found in the secondary references which use free radically curable systems with their matrix (such as acrylates) as does Korishima et al. JP 05-046061. **The closest prior art is Korishima et al. JP 05-046061, not Dhar et al.** Further, it is clear from

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Korishima et al. JP 05-046061 that the sensitivity will be different from other photoinitiation systems, as its sensitivity in the red and infrared is heightened. [0003-0004,0023]. Therefore the data asserted by the applicant's representative as unobvious is not. The rejection stands.

4. Claims 1-4,6-9,14 and 16-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Korishima et al. JP 05-046061, in view of Hegel et al. '008.

Hegel et al. '008 teach in example 1, an acrylate monomer and IRG-784 as the photoinitiator mixed with matrix precursors dibutyltin diacetate, diisocyanate terminated polypropylene glycol and dihydroxypolypropylene glycol, which are placed between 1.2 mm substrates with a 500 micron spacer and after curing of the matrix [0031-0040]. The formation of holograms using these is disclosed. [0052-0056]. Useful photoactive monomers including acrylates are disclosed as useful in this system. [0029]. A reduction in shrinkages of the hologram is also realized. The provision of antireflection coatings on one or both of the substrates is disclosed. [0016,0018]. The substrates may be 0.5-1.3 mm thick [0019].

It would have been obvious to one skilled in the art to modify the cited example of Korishima et al. JP 05-046061 by adding the components to form an insitu matrix and place the composition between two glass slides with a spacer of 500 microns to allow a thicker hologram to be formed and reduce shrinkage as discussed by Hegel et al. '008.

The rejection stands for the reasons above as no further arguments were directed at this rejection.

5. Claims 1-9,14 and 16-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Korishima et al. JP 05-046061, in view of Dhar et al. '104.

Dhar et al. '104 teach in example 1, an acrylate monomer and CGI-784 as the photoinitiator mixed with matrix precursors dibutyltin dilaurate, diisocyanate terminated polypropylene glycol and dihydroxypolypropylene glycol [0091-0092], placed between two PMMA discs with a 750 microns spacers and the matrix cured [0096-0101]. The ability to form thick holographic recording layers of more than 200 microns is disclosed [0072]. Useful photoactive monomers including acrylates are disclosed as useful in this system [0030]. A reduction in shrinkage of holograms is also realized (7/1-48). The use of various reactions including hydrosilation and the like to form the matrix is disclosed.. The matrix can also be made form polyols which are fluorinated [0027]

It would have been obvious to one skilled in the art to modify the cited example of Korishima et al. JP 05-046061 by adding the components to form an insitu matrix and place the composition between two glass slides with a spacer of 200-500 microns to allow a thicker hologram to be formed and reduce shrinkage as discussed by Dhar et al. '104.

The rejection stands for the reasons above as no further arguments were directed at this rejection.

6. Claims 1-4,6-9,14 and 16-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Korishima et al. JP 05-046061, in view of Dhar et al. '551, further in view of Gottschalk et al. '541 or Adair et al. '414.

Gottschalk et al '541 teach the use of dye borate complexes to cure free radical polymerizable systems. The cationic dyes may be methine, polymethine, triarylmethane, indolene, thiazine, xanthene, oxazine, acridine, cyanine, carbocyanine, hemicyanine, rhodamine and azomethine dyes. (5/25-6/54). The initiation is described as more efficient because of the

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close association of the dye, which absorbs the light, and the borate anion, which generates the free radical and facilitates the transfer of the energy from the excited dye to the borate. (3/4-14).

Adair et al. '414 teach the use of cationic transition metal complexes with borate anions to cure free radically polymerizable systems, where the ligands on the complexes can be pyridine, bipyridines, phenanthrolines (the latter have two chelation sites) and the like (4/37-6/49)/

To address the other embodiments bounded by the claims where the dye cations are not cyanine, the examiner holds that it would have been obvious to one skilled in the art to modify the composition rendered obvious by the combination of Korishima et al. JP 05-046061, and Dhar et al. '551 by using other known dye/borate photoinitiators such as those taught by Gottschalk et al. '541 or Adair et al. '414 with a reasonable expectation of the free radical components in the resulting combination being photocurable based upon the common usage of free radically curable monomers in all the references.

The applicant repeats the argument of the ammonium cation, which has been addressed above. The applicant also argues that the dyes of Adair et al. '414 are not within formula 2. The examiner points to fact that the ligands are not limited to those illustrated in section of the [0075] of the instant specification. The applicant could exclude the ligands of Adair et al. '414, but has chosen not to do so, so the position argued is flawed. Further based upon the fact that Gottschalk et al. teaches cyanine dye/ borate initiators, which embraces those of Korishima et al. JP 05-046061 and uses them for the same purpose (photoinitiation), the argument that they are not combinable by the applicant are unconvincing.

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7. Claims 1-4,6-9,14 and 16-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Korishima et al. JP 05-046061, in view of Hegel et al. '008, further in view of Gottschalk et al. '541 or Adair et al. '414.

To address the other embodiments bounded by the claims where the dye cations are not cyanine, the examiner holds that it would have been obvious to one skilled in the art to modify the composition rendered obvious by the combination of Korishima et al. JP 05-046061, and Hegel et al. '008 by using other known dye/borate photoinitiators such as those taught by Gottschalk et al. '541 or Adair et al. '414 with a reasonable expectation of the free radical components in the resulting combination being photocurable based upon the common usage of free radically curable monomers in all the references.

The rejection stands for the reasons above as no further argument were directed at this line of rejection.

8. Claims 1-12,14 and 16-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Korishima et al. JP 05-046061, in view of Dhar et al. '104, further in view of Gottschalk et al. '541 or Adair et al. '414.

To address the other embodiments bounded by the claims where the dye cations are not cyanine, the examiner holds that it would have been obvious to one skilled in the art to modify the composition rendered obvious by the combination of Korishima et al. JP 05-046061, and Dhar et al. '104 by using other known dye/borate photoinitiators such as those taught by Gottschalk et al. '541 or Adair et al. '414 with a reasonable expectation of the free radical components in the resulting combination being photocurable based upon the common usage of free radically curable monomers in all the references.

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The rejection stands for the reasons above as no further argument were directed at this line of rejection.

9. Claims 1-4,6-10, and 12-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Korishima et al. JP 05-046061, in view of Hegel et al. '008, further in view of Gottschalk et al. '541 or Adair et al. '414, further in view of Horimai et al., WO 02/15176.

Horimai et al., WO 02/15176 (Hormai et al. '891 is US equivalent) teaches with respect to figure 1, a holographic recording medium which comprises a substrate (2), a holographic recording layer (photopolymers) (3), a second substrate (4) and a reflective layer (5). The reflective layer and the recording layer can be next to each other (12/5-23; 11/40-64). The interference fringes results from the interferences from the light passing through the layer toward the reflective layer and that reflected back into the laser from the reflective layer. (col 5. ?; 4/53-5/7).

In addition to the basis provided above, it would have been obvious to one skilled in the art to modify the embodiments rendered obvious by the combination of Korishima et al. JP 05-046061, with Hegel et al. '008 and either of (Gottschalk et al. '541 or Adair et al. '414) as set forth above by adding a reflective layer on the further substrate as taught Horimai et al., WO 02/15176 to allow holographic recording without a second beam.

No arguments were directed at this rejection.

10. Claims 1-10, and 12-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Korishima et al. JP 05-046061, in view of Dhar et al. '104, further in view of Gottschalk et al. '541 or Adair et al. '414, further in view of Horimai et al., WO 02/15176.

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In addition to the basis provided above, it would have been obvious to one skilled in the art to modify the embodiments rendered obvious by the combination of Korishima et al. JP 05-046061, with Dhar et al. '104 and either of (Gottschalk et al. '541 or Adair et al. '414) as set forth above by adding a reflective layer on the further substrate as taught Horimai et al., WO 02/15176 to allow holographic recording without a second beam.

No arguments were directed at this rejection.

11. The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). See, e.g., *In re Berg*, 140 F.3d 1428, 46 USPQ2d 1226 (Fed. Cir. 1998); *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) or 1.321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

12. Claims 1-4,8-10 and 12-20 are provisionally rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-16 of copending Application No. 11/201,815 (US 2006/0040185) in view of Korishima et al. JP 05-046061.

It would have been obvious to one skilled in the art to modify the claims of the copending application 11/201,815, by using photopolymerizable compounds and initiators known to be

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useful with them in the formation of holographic patterns, such as those disclosed by Korishima et al. JP 05-046061 with a reasonable expectation of forming a useful holographic recording medium.

This is a provisional obviousness-type double patenting rejection.

The applicant stated that a terminal disclaimer might be filed at a later date.

13. **THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

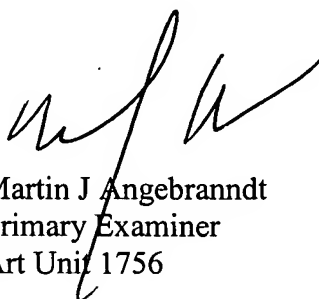
A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

14. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Martin J. Angebrannndt whose telephone number is 571-272-1378. The examiner can normally be reached on Monday-Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Mark Huff can be reached on 571-272-1385. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.



Martin J. Angebranndt
Primary Examiner
Art Unit 1756

5/22/2007